Application No. 10/762,616 Amdt. dated 12 June 2009 Reply to Office Action of 4 February 2009 and the Advisory Action of 1 June 2009

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims 1-22:

- 1. (currently amended) A pharmaceutical composition for oral administration including tinidazole and fluconazole or a stereoisomer or a stereoisomeric <u>uniform</u> mixture thereof, comprising from about 50 to less than 150 mg fluconazole and from about 1000 to less than 2000 mg tinidazole in a unit dose.
- 2. (currently amended) A pharmaceutical composition characterized by secnidazole and fluconazole or a stereoisomer or a stereoisomeric mixture thereof, comprising a unit dosage having a uniform mixture consisting essentially of from about 50 to less than 150 mg of fluconazole and from about 1000 to less than 2000 mg of secnidazole.
- 3. (previously presented) The pharmaceutical composition according to Claim 2, characterized by a percent by weight of 75% \pm 20% secnidazole and a percent by weight of 6% \pm 2% fluconazole.
 - 4. (cancelled)
- 5. (previously presented) The pharmaceutical composition according to Claim 3, characterized by a tablet form.
- 6. (previously presented) The pharmaceutical composition according to Claim 2, characterized by a vehicle mixture of acceptable pharmaceutical vehicles that comprises microcrystalline cellulose, sodium glycolate of starch, polyvinylpyrrolidone, magnesium stearate and white opadry.

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(cancelled)
(cancelled)
(cancelled)
(cancelled)
(cancelled)
(cancelled)

(cancelled)

12.

- 13. (previously presented) The pharmaceutical composition according to Claim 1, characterized by a percent by weight of 75% \pm 20% tinidazole and a percent by weight of 6% \pm 2% fluconazole.
 - 14. (cancelled)
- 15. (previously presented) The pharmaceutical composition according to Claim 13, characterized by a tablet form.
- 16. (previously presented) The pharmaceutical composition according to Claim 1, characterized by a vehicle mixture of acceptable pharmaceutical vehicles that comprises microcrystalline cellulose, sodium glycolate of starch, polyvinylpyrrolidone, magnesium stearate and white opadry.

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- 17. (previously presented) The pharmaceutical composition according to Claim 1, characterized by a percent by weight of 75% tinidazole.
- 18. (previously presented) The pharmaceutical composition according to Claim 2, characterized by a percent by weight of 75% secnidazole.
- 19. (previously presented) The pharmaceutical composition according to Claim 1, characterized by a percent by weight of 5.7% fluconazole.
- 20. (previously presented) The pharmaceutical composition according to Claim 2, characterized by a percent by weight of 5.7% fluconazole.
- 21. (previously presented) The pharmaceutical composition according to Claim 1, wherein the composition is characterized by about 112.5 mg of fluconazole and about 1.5 g of tinidazole.
- 22. (previously presented) The pharmaceutical composition according to Claim 2, wherein the composition is characterized by about 112.5 mg of fluconazole and about 1.5 g of secnidazole.